



## **DEPARTMENT OF JUSTICE**

### **Drug Enforcement Administration**

**[Docket No. DEA-392]**

#### **Importer of Controlled Substances Application: Galephar Pharmaceutical Research Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on June 10, 2019, Galephar Pharmaceutical Research Inc., #100 Carr 198 Industrial Park, Juncos, Puerto Rico, 00777 applied to be registered as an importer of the following basic class of controlled substance:

<b>Controlled Substance</b>	<b>Drug Code</b>	<b>Schedule</b>
Hydromorphone	9150	II

The company plans to import the listed controlled substance in finished dosage form for analytical purpose only.

Dated: August 22, 2019.

Neil D. Doherty,  
*Acting Assistant Administrator.*

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